In the claims:

1. (currently amended) A compound of formula (I)

$$R^5$$
 R^4
 R^3
 R^2
 R^7
 R^1

X is CH2 or SO2

R1 is an optionally substituted aryl or heteroaryl ring;

R² is carboxy, cyano, -C(O)CH₂OH, -CONHR⁸, -SO₂NHR⁹, tetrazol-5-yl, or SO₃H, or a group of formula (VI)

(IV)

where R⁸ is selected from hydrogen, alkyl, aryl, cyano, hydroxy, -SO₂R¹² where R¹² is alkyl, aryl, heteroaryl, or haloalkyl, or R⁸ is a group-(CHR¹³)_i-COOH where r is an integer of 1-3 and each R¹³ group is independently selected from hydrogen or alkyl; R⁹ is hydrogen, alkyl, optionally substituted aryl such as optionally substituted phenyl or optionally subtituted substituted heteroaryl such as 5 or 6 membered heteroaryl groups, or a group COR¹⁴ where R¹⁴ is alkyl, aryl, heteroaryl or haloalkyl; R¹⁰-and R¹¹ are independently selected from hydrogen or alkyl, particularly C_{1,4}-alkyl;

R³ is hydrogen, a functional group, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted aryl, optionally substituted heterocyclyl,

optionally substituted alkoxy, optionally substituted aralkyl, optionally substituted aralkyloxy; or optionally substituted cycloalkyl;

R⁴ is a group NHCOR¹⁵; or NHSO₂R¹⁵ or OCONR¹⁶R¹⁷ where R¹⁵ is optionally substituted alkyl, optionally substituted aryl or optionally substituted heteroaryl-and R¹⁶ and R¹⁷ are independently substituted from hydrogen, optionally substituted alkyl, optionally substituted aryl and optionally substituted heteroaryl, with the provise that at least one of R¹⁶ or R¹⁷ is other than hydrogen, or R¹⁶ and R¹⁷ together with the nitrogen atom to which they are attached form an optionally substituted heterocyclic ring which optionally contains further heteroatoms; and R⁵, R⁶ and R⁷ are independently selected from hydrogen, a functional group or an optionally substituted hydrocarbyl group groups or optionally substituted heterocyclic groups; and further provided that when R⁴ is a group NHCOR¹⁵, R¹⁵ is substituted alkyl, optionally substituted aryl or optionally substituted heteroaryl.

- 2. (currently amended) A compound according to claim 1 wherein a group R¹⁵, R¹⁶ and R¹⁷ as they appear as it appears in the definition of R⁴, is substituted by at least one functional group, or an aryl or heterocyclyl group groups, either of which may themselves be substituted by one or more functional groups or further aryl or heterocyclyl groups.
- 3. (currently amended) A compound according to any one of the preceding claims claim 1 wherein R⁴ is a group NHCOR¹⁵ or NHSO₂R¹⁵ and R¹⁵ is a substituted alkyl group or an optionally substituted heterocyclyl substituted heterocyclyl or optionally substituted phenyl group.
- 4. (currently amended) A compound according to claim 3 wherein R^{15} is alkyl substituted by a group of formula $NR^{19}R^{20}$ where R^{19} and R^{20} are independently selected from hydrogen or optionally substituted hydrocarbyl, or R^{19} and R^{20} together form an optionally substituted ring which optionally contains further heteroatoms such as $S(O)_m$, oxygen and nitrogen, n is an integer of 1 or 2, and m is 1 or 2.
- 5. (currently amended) A compound according to any one of the preceding claims claim 1 where R² is carboxy.

- 6. (currently amended) A compound according to any one of the preceding claims claim 1 wherein R¹ is 3,4-dichlorophenyl, 3-fluoro-4-chlorophenyl, 3-chloro-4-fluorophenyl or 2,3-dichloropyrid-5-yl.
- 7. (currently amended) A compound according to any one of the preceding claims claim 1 where X is CH₂.
- 8. (currently amended) A process for preparing a compound according to claim 1 which process comprises either

(a) where R4 is NHCOR45 or NHSO2R45, reacting a compound of formula (VII)

where X, R¹, R³, R⁵, R⁶ and R⁷ are as defined in claim 1, and R² is a group R² as defined in relation to formula (I) or a protected form thereof, with a compound of formula (VIII)

(VIII)

where Z is a leaving group and R^{22} is a group COR^{15} or SO_2R^{15} where R^{15} is group R^{15} as defined in relation to formula (I) or a precursor thereof;

or (b) where R4 is a group OCONR 16R17, reacting a compound of formula (VIIA)

(VIIA)

where X, R^{2'}, R¹, R², R⁵, R⁶ and R⁷ are as defined claim 1 and R² is a group R² as defined in claim 1 or a protected form thereof, with a compound of formula (VIIIA)

Z-CONR¹⁶R¹⁷ (VIIIA)

where Z, R¹⁶ and R¹⁷ are as defined above; and thereafter if desired or necessary:

- (i) converting a precursor group R^{15} to a group R^{15} and/or converting a group R^{15} to a different such group; and
- (ii) deprotecting a group R2' to a group R2.
- 9. (currently amended) A pharmaceutical composition comprising a compound according to any one of claims 1 to 7 claim 1 in combination with a pharmaceutically acceptable carrier.
- 10. (currently amended) A method for antagonizing an MCP-1 (Monocyte Chemoattractant Protein-1) or RANTES (Regulated upon Activation, Normal T-cell Expressed and Secreted) mediated effect in a warm blooded animal in need of such treatment comprising administering to said animal an effective amount of aA compound according to any one of claims-1 to 7claim 1, a pharmaceutically acceptable salt, or an in vivo hydrolysable ester thereof, for use in the preparation of a medicament for use in the treatment of disease mediated by monocyto chemoattractant protein 1 or RANTES (Regulated upon Activation, Normal T cell Expressed and Secreted), such as inflammatory disease.
- 11. (new) A method for treating inflammation in a warm blooded animal in need of such treatment comprising administering to said animal an effective amount of a compound according to claim 1, a pharmaceutically acceptable salt, or an *in vivo* hydrolysable ester thereof.